

**AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

Claims 1-27 (Cancelled).

Claim 28 (Previously presented): A method for treating cerebral ischemia in a mammal comprising peripherally administering to said mammal in need thereof a non-toxic amount of erythropoietin effective to exert a neuroprotective effect.

Claim 29 (Previously presented): The method of Claim 28 wherein said administering is carried out in a vascular fashion.

Claim 30 (Previously presented): The method of Claim 29 wherein said vascular administration is intravenous.

Claim 31 (Previously presented): The method of Claim 28, 29, or 30 wherein said erythropoietin is administered for the treatment of stroke.

Claim 32 (Previously presented): The method of Claim 28, 29 or 30 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.

Claim 33 (Previously presented): The method of Claim 28, 29, or 30 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin.

Claim 34 (Previously presented): A method for treating cerebral ischemia in a human subject comprising peripherally administering to said human subject in need thereof a non-toxic amount of erythropoietin effective to exert a neuroprotective effect.

Claim 35 (Previously presented): The method of Claim 34 wherein said administering is carried out in a vascular fashion.

Claim 36 (Previously presented): The method of Claim 34 wherein said vascular administration is intravenous.

Claim 37 (Previously presented): The method of Claim 34, 35, or 36 wherein said erythropoietin is administered for the treatment of stroke.

Claim 38 (Previously presented): The method of Claim 34, 35, or 36 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.

Claim 39 (Previously presented): The method of Claim 34, 35, or 36 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin.

Claims 40-51 (Cancelled).

Claim 52 (Previously presented): A method for treating cerebral ischemia in a mammal comprising peripherally administering to said mammal in need thereof an amount of erythropoietin effective to exert a neuroprotective effect without an increase in hematocrit in said mammal.

Claim 53 (Previously presented): The method of Claim 52 wherein said administering is carried out in a vascular fashion.

Claim 54 (Previously presented): The method of Claim 53 wherein said vascular administration is intravenous.

Claim 55 (Previously presented): The method of Claim 52, 53, or 54 wherein said erythropoietin is administered for the treatment of stroke.

Claim 56 (Previously presented): The method of Claim 52, 53, or 54 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.

Claim 57 (Previously presented): The method of Claim 52, 53, or 54 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin.

Claim 58 (Previously presented): A method for treating cerebral ischemia in a human subject comprising peripherally administering to said human subject in need thereof an amount of erythropoietin effective to exert a neuroprotective effect without an increase in hematocrit in said human subject.

Claim 59 (Previously presented): The method of Claim 58 wherein said administering is carried out in a vascular fashion.

Claim 60 (Previously presented): The method of Claim 58 wherein said vascular administration is intravenous.

Claim 61 (Previously presented): The method of Claim 58, 59, or 60 wherein said erythropoietin is administered for the treatment of stroke.

Claim 62 (Previously presented): The method of Claim 58, 59, or 60 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.

Claim 63 (Previously presented): The method of Claim 58, 59, or 60 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin.

Claim 64 (Previously presented): The method of Claim 28, 29, 30, 34, 35, 36, 52, 53, 54, 58, 59, or 60 wherein said erythropoietin is an erythropoietin derivative.